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COMPLETE LISTING OF ALL CLAIMS, WITH MARKINGS AND STATUS IDENTIFIERS

(Currently amended claims showing deletions by $\frac{by-strikethrough}{underlining}$ and additions by $\frac{underlining}{}$

1 - 31 (canceled)

32 (currently amended): A pharmaceutical composition for the treatment of hyperlipidemia in a patient in need thereof, comprising a therapeutically effective amount of a somatostatin type 5 receptor selective an agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor and a binding affinity (Ki) of less than 5nM for the somatostatin type-5 receptor, wherein said therapeutically effective amount is an amount that is effective for the treatment of hyperlipidemia in said patient.

33 (canceled)

34 (currently amended): A pharmaceutical composition according to claim 32, wherein said somatostatin type 5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki of less than 2 nM for the somatostatin type-5 receptor.

35 (currently amended): A pharmaceutical composition according to claim 32, wherein said somatostatin type 5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has

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a Ki for the type-5 somatostatin receptor that is at least 10 times less than its Ki for the somatostatin type-2 receptor.

36 - 37 (canceled)

38 (currently amended): A pharmaceutical composition for lowering the amount of triacylglycerols in the blood of a patient in need of such lowering, comprising a therapeutically effective amount of a somatostatin type 5 agonist receptor selective an selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor and a binding affinity (Ki) of less than 5nM for the somatostatin type-5 receptor, wherein said therapeutically effective amount is an amount that is effective for lowering the amount of triacylglycerols in the blood of said patient.

39 (canceled)

40 (currently amended): A pharmaceutical composition according to claim 38, wherein said somatostatin type 5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki of less than 2 nM for the somatostatin type-5 receptor.

41 (currently amended): A pharmaceutical composition according to claim 38, wherein said somatostatin type-5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the

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somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki for the type-5 somatostatin receptor that is at least 10 times less than its Ki for the somatostatin type-2 receptor.

42 - 43 (canceled)

44 (currently amended): A pharmaceutical composition for lowering the amount of glycerol in the blood of a in need of such patient lowering, comprising therapeutically effective amount of a somatostatin type 5 receptor selective <u>an</u> agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor and a binding affinity (Ki) of less than 5nM for somatostatin type-5 receptor, wherein said therapeutically effective amount is an amount that is effective for lowering the amount of glycerol in the blood of said patient.

45 (canceled)

46 (currently amended): A pharmaceutical composition according to claim 44, wherein said somatostatin type-5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki of less than 2 nM for the somatostatin type-5 receptor.

47 (currently amended): A pharmaceutical composition according to claim 44, wherein said somatostatin type 5

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receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki for the type-5 somatostatin receptor that is at least 10 times less than its Ki for the somatostatin type-2 receptor.

48 - 49 (canceled)

(currently amended): A pharmaceutical composition for lowering the amount of cholesterol in the blood of a need of patient in such lowering, comprising therapeutically effective amount of a somatostatin type 5 receptor selective selective for the an agonist somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor and a binding affinity (Ki) of less than 5nM for somatostatin type-5 receptor, wherein therapeutically effective amount is an amount that effective for lowering the amount of cholesterol in the blood of said patient.

51 (canceled)

52 (currently amended): A pharmaceutical composition according to claim 50, wherein said somatostatin type 5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either the somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki of less than 2 nM for the somatostatin type-5 receptor.

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53 (currently amended): A pharmaceutical composition according to claim 50, wherein said somatostatin type 5 receptor agonist selective for the somatostatin type-5 receptor and having a higher binding affinity for the somatostatin type-5 receptor than for either somatostatin type-1, type-2, type-3 or type-4 receptor has a Ki for the type-5 somatostatin receptor that is at least 10 times less than its Ki for the somatostatin type-2 receptor.

54 - 55 (canceled)